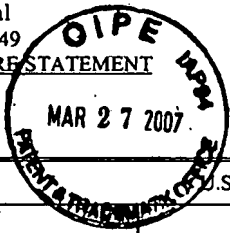


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EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	REFERENCE PROVIDED*
/L.H./	AA1	5,413,996	05/09/1995	Bodor	not required, per 69 Fed. Reg. 56481
	AB1	5,670,497	09/23/1997	Bold <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AC1	5,750,343	05/12/1998	Maag <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AD1	5,750,493	05/12/1998	Sommadossi <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AE1	5,874,577	02/23/1999	Chen <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AF1	5,914,332	06/22/1999	Sham <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AG1	6,072,053	06/06/2000	Vince <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AH1	6,312,662	11/06/2001	Erion <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AI1	6,319,946	11/20/2001	Hale <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AJ1	6,767,900	07/27/2004	Ubasawa <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AK1	2001/031773	10/18/2001	Camden	not required, per 69 Fed. Reg. 56481
	AL1	2002/0119443 A1	08/29/2002	Becker <i>et al.</i>	not required, per 69 Fed. Reg. 56481
	AM1	2003/109498	06/12/2003	Yuasa <i>et al.</i>	not required, per 69 Fed. Reg. 56481
↓	AN1	2004/0121316 A1	06/24/2004	Birkus <i>et al.</i>	not required, per 69 Fed. Reg. 56481

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EXAMINER INITIAL		DOCUMENT NUMBER	DATE	COUNTRY	REFERENCE PROVIDED*	TRANSLATION
/L.H./	AO1	0 267 050 A2	05/11/1988	Europe		Yes No
	AP1	0 441 192 A2	08/14/1991	Europe		x (abstract only) Yes No
	AQ1	0 465 297 A1	01/08/1992	Europe		Yes No
	AR1	0 531 597 A1	03/17/1993	Europe		Yes No
	AS1	0 632 048 A1	01/04/1995	Europe		Yes No
	AT1	0 786 455 A1	07/30/1997	Europe		Yes No
	AU1	0 852 233 A1	07/08/1998	Europe		Yes No
	AV1	0 919 562 A1	06/02/1999	Europe		Yes No
↓	AW1	1 295 879 A1	03/26/2003	Europe		Yes No


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/L.H./	AX1	WO 88/06158	08/25/1988	PCT		x (abstract only)	Yes No		
	AY1	WO 91/19721	12/26/1991	PCT			Yes No		
	AZ1	WO 92/00988	01/23/1992	PCT			Yes No		
	AA2	WO 92/18520	10/29/1992	PCT		x (abstract only)	Yes No		
	AB2	WO 93/12123	06/24/1993	PCT			Yes No		
	AC2	WO 93/24510	12/09/1993	PCT		x (Abstract only)	Yes No		
	AD2	WO 96/14314	05/17/1996	PCT			Yes No		
	AE2	WO 96/40156	12/19/1996	PCT			Yes No		
	AF2	WO 97/15588 A1	05/01/1997	PCT			Yes No		
	AG2	WO 98/04569	02/05/1998	PCT			Yes No		
	AH2	WO 98/11906	03/26/1998	PCT			Yes No		
	AI2	WO 99/62921	12/09/1999	PCT			Yes No		
	AJ2	WO 00/04033	01/27/2000	PCT			Yes No		
	AK2	WO 00/52015	08/09/2000	PCT			Yes No		
	AL2	WO 01/13957 A2	03/01/2001	PCT			Yes No		
	AM2	WO 01/13957 A3	03/01/2001	PCT			Yes No		
	AN2	WO 01/17982 A1	03/15/2001	PCT			Yes No		
	AO2	WO 01/19320 A2, A3	03/22/2001	PCT			Yes No		
	AP2	WO 01/39724 A2	06/07/2001	PCT			Yes No		
	AQ2	WO 01/39724 A3	10/18/2001	PCT			Yes No		
V	AR2	WO 01/46204 A1	06/28/2001	PCT			Yes No		
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/L.H./	AS2	WO 01/64693 A1	09/07/2001	PCT		x (Abstract only)	Yes No
	AT2	WO 01/96329 A1	12/20/2001	PCT		x (Abstract only)	Yes No
	AU2	WO 01/96354 A1	12/20/2001	PCT		x (Abstract only)	Yes No
	AV2	WO 02/03997 A1	01/17/2002	PCT			Yes No
	AW2	WO 02/06292 A1	01/24/2002	PCT			Yes No
	AX2	WO 02/08241 A2	01/31/2002	PCT			Yes No
	AY2	WO 02/103008 A2	12/27/2002	PCT			Yes No
	AZ2	WO 02/103008 A3	11/27/2003	PCT			Yes No
	AA3	WO 02/14344 A2	02/21/2002	PCT			Yes No
	AB3	WO 02/057425 A2	07/25/2002	PCT			Yes No
	AC3	WO 02/100415 A2, A3	12/19/2002	PCT			Yes No
	AD3	WO 03/028737 A1	04/10/2003	PCT			Yes No
	AE3	WO 03/050129 A1	06/19/2003	PCT			Yes No
	AF3	WO 03/059255 A2	07/24/2003	PCT			Yes No
	AG3	WO 03/064383 A2	08/07/2003	PCT			Yes No
	AH3	WO 03/066005 A2	08/14/2003	PCT			Yes No
	AI3	WO 03/080078 A1	10/02/2003	PCT			Yes No
	AJ3	WO 03/090690 A2	11/06/2003	PCT			Yes No
	AK3	WO 04/096234 A2	11/11/2004	PCT			Yes No
V	AL3	WO 05/011709 A1	02/10/2005	PCT			Yes No
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/L.H./ 	AM3	Abdel-Meguid <i>et al.</i> , "Inhibition of Human Immunodeficiency Virus-1 Protease by a C ₂ -Symmetric Phosphinate. Synthesis and Crystallographic Analysis", <i>Biochemistry</i> , 32(31):7972-7980 (1993)			
	AN3	Allen <i>et al.</i> , "CI-1040 (PDI84352), a Targeted Signal Transduction Inhibitor of MEK (MAPKK)", <i>Seminars in Oncology</i> , 30(5, Suppl. 16):105-116 (2003)			
	AO3	Andrade <i>et al.</i> , "HIV-Related Drug Metabolism and Cytochrome P450 Enzymes" <i>AIDS Clinical Care</i> , 12(11):91-95 (2000)			
	AP3	Ballatore <i>et al.</i> , "Synthesis and Evaluation of Novel Amidate Prodrugs of PMEA and PMPA", <i>Bioorganic & Medicinal Chemistry Letters</i> , 11:1053-1056 (2001)			
	AQ3	Bantia <i>et al.</i> , "Purine Nucleoside Phosphorylase Inhibitor BCX-1777 (Immucillin-H)—A Novel Potent and Orally Active Immunosuppressive Agent", <i>International Immunopharmacology</i> , 1:1199-1210 (2001)			
	AR3	Beauchamp, <i>et al.</i> , "Guanine, Pyrazolo[3,4-d]pyrimidine, and Triazolo[4,5-d]pyrimidine(8-Azaguanine) Phosphonate Acyclic Derivatives as Inhibitors of Purine Nucleoside Phosphorylase", <i>Journal of Medicinal Chemistry</i> , 39:949-956 (1996)			
	AS3	Borhani <i>et al.</i> , "A-420983: A Potent, Orally Active Inhibitor of lck with Efficacy in a Model of Transplant Rejection", <i>Bioorganic & Medicinal Chemistry Letters</i> , 14:2613-2616 (2004)			
	AT3	Bzowska <i>et al.</i> , "Purine Nucleoside Phosphorylases: Properties, Functions, and Clinical Aspects", <i>Pharmacology & Therapeutics</i> , 88:349-425 (2000)			
	AU3	Chapman <i>et al.</i> , "Practical Synthesis, Separation, and Stereochemical Assignment of the PMPA Pro-Drug GS-7340", <i>Nucleosides, Nucleotides & Nucleic Acids</i> , 20(4-7):621-628 (2001)			
AV3	Charvet <i>et al.</i> , "Inhibition of Human Immunodeficiency Virus Type I Replication by Phosphonoformate- and Phosphonoacetate-2',3'-Dideoxy-3'-thiacytidine Conjugates", <i>Journal of Medicinal Chemistry</i> , 37(14):2216-2223 (1994)				
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/L.H./	AW3	Clark <i>et al.</i> , Abstract, "Mycophenolic Acid Analogues as Potential Agents Against West Nile Virus Infection", Institute for Antiviral Research, Utah State University			
	AX3	Conklyn <i>et al.</i> , "The JAK3 Inhibitor CP-690550 Selectively Reduces NK and CD8+ Cell Numbers in Cynomolgus Monkey Blood Following Chronic Oral Dosing", <i>Journal of Leukocyte Biology</i> , 76:1-8 (2004)			
	AY3	De Clercq, "Chemotherapy of Human Immunodeficiency Virus (HIV) Infection: Anti-HIV Agents Targeted at Early Stages in the Virus Replicative Cycle", <i>Biomedicine and Pharmacotherapy</i> , 50(5):207-215 (1996)			
	AZ3	De Clercq, "Highlights in the Development of New Antiviral Agents", <i>Mini Reviews in Medicinal Chemistry</i> , 2(2):163-175 (2002)			
	AA4	De Clercq, "New Developments in Anti-HIV Chemotherapy", <i>Current Medicinal Chemistry</i> , 8(13):1543-1572 (2001)			
	AB4	Dvořáková <i>et al.</i> , "Synthesis of 2'-Aminomethyl Derivatives of N-(2-(Phosphonomethoxy)ethyl) Nucleotide Analogues as Potential Antiviral Agents", <i>J. Med. Chem.</i> , 39(17):3263-3268 (1996)			
	AC4	Evans <i>et al.</i> , "Exploring Structure-Activity Relationships of Transition State Analogues of Human Purine Nucleoside Phosphorylase", <i>J. Med. Chem.</i> , 46(15):3412-3423 (2003)			
	AD4	Gobec <i>et al.</i> , Phosphonate Inhibitors of Antigen 85C, A Crucial Enzyme Involved in the Biosynthesis of the <i>Mycobacterium Tuberculosis</i> Cell Wall", <i>Bioorganic and Medicinal Chemistry Letters</i> , 14:3559-3562 (2004)			
	AE4	Gorin <i>et al.</i> , "A Novel Esterification Procedure Applied to Synthesis of Biologically Active Esters of Foscarnet", <i>Tetrahedron Letters</i> , 38(16):2791-2794 (1997)			
↓	AF4	Gumina <i>et al.</i> , "Advances in antiviral agents for Hepatitis B Virus", <i>Antiviral Chemistry & Chemotherapy</i> , 12(Suppl. 1):93-117 (2001)			
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/L.H./	AG4	Hakimelahi <i>et al.</i> , "Design, Synthesis, and Structure-Activity Relationship of Novel Dinucleotide Analogs as Agents Against Herpes and Human Immunodeficiency Viruses", <i>Journal of Medicinal Chemistry</i> , 38(23):4648-4659 (1995)	
	AH4	Hammond <i>et al.</i> , "Alkylglycerol Prodrugs of Phosphonoformate are Potent In Vitro Inhibitors of Nucleoside-Resistant Human Immunodeficiency Virus Type 1 and Select for Resistance Mutations that Suppress Zidovudine Resistance", <i>Antimicrobial Agents and Chemotherapy</i> , 45(6):1621-1628 (2001)	
	AI4	Hegedus <i>et al.</i> , "Synthesis of 4'-Methyl and 4'-Cyano Carbocyclic 2',3'-Didehydro Nucleoside Analogues via 1,4-Addition to Substituted Cyclopentenones", <i>J. Org. Chem.</i> , 69(24):8492-8495 (2004)	
	AJ4	Herczegh <i>et al.</i> , "Osteoadsorptive Bisphosphonate Derivatives of Fluoroquinolone Antibacterials", <i>J. Med. Chem.</i> , 45:2338-2341 (2002)	
	AK4	Hirabayashi <i>et al.</i> , "Bone-Specific Drug Delivery Systems: Approaches via Chemical Modification of Bone-Seeking Agents", <i>Clinical Pharmacokinetics</i> , 42(15):1319-1330 (2003)	
	AL4	Holý <i>et al.</i> , "Synthesis of N-(2-Phosphonylmethoxyethyl Derivatives of Heterocyclic Bases", <i>Collect. Czech. Chem. Commun.</i> , 54:2190-2210 (1989)	
	AM4	International Search Report for International Application No. PCT/EP2003/012423, mailed February 11, 2005	
	AN4	International Search Report for International Application No. PCT/US2004/035083, mailed August 5, 2005	
	AO4	International Search Report for International Application No. PCT/US2004/035084, mailed March 2, 2005	
	AP4	International Search Report for International Application No. PCT/US2004/035085, mailed March 2, 2005	
↓	AQ4	Jain <i>et al.</i> , "Characterization of Pharmacological Efficacy of VX-148, a New, Potent Immunosuppressive Inosine 5'-Monophosphate Dehydrogenase Inhibitor", <i>Journal of Pharmacology and Experimental Therapeutics</i> , 302(3):1272-1277 (2002)	
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/L.H./	AR4	Karpenko <i>et al.</i> , "Synthesis and Antiherpetic Activity of Acyclovir Phosphonates", <i>Nucleosides, Nucleotides & Nucleic Acids</i> , 22(3):319-328 (2003)			
	AS4	Kato <i>et al.</i> , "Enantio- and diastereoselective Synthesis of 4' - α -Substituted Carbocyclic Nucleosides", <i>Tetrahedron: Asymmetry</i> , 9:911-914 (1998)			
	AT4	Kato <i>et al.</i> , "Stereoselective Synthesis of 4' - α -alkylcarbovir Derivatives Based on an Asymmetric Synthesis or Chemo-Enzymatic Procedure", <i>Chemical & Pharmaceutical Bulletin</i> , 47(9):1256-1264 (1999)			
	AU4	Kilpatrick <i>et al.</i> , "Intravenous and Oral Pharmacokinetic Study of BCX-1777, a Novel Purine Nucleoside Phosphorylase Transition-State Inhibitor, <i>In vivo</i> Effects on Blood 2'-Deoxyguanosine in Primates", <i>International Immunopharmacology</i> , 3:541-548 (2003)			
	AV4	Kim <i>et al.</i> , "Regiospecific and Highly Stereoselective Electrophilic Addition to Furanoid Glycols: Synthesis of Phosphonate Nucleotide Analogues with Potent Activity Against HIV", <i>J. Org. Chem.</i> , 56(8):2642-2647 (1991)			
	AW4	Kinsky <i>et al.</i> , "Inhibition of Cell Proliferation by Putative Metabolites and Non-Degradable Analogs of Methotrexate- γ -Dimyristoylphosphatidylethanolamine", <i>Biochimica et Biophysica Acta</i> , 917(2):211-218 (1987)			
	AX4	Kinsky <i>et al.</i> , "Effect of Liposomes Scentitized with Methotrexate- γ -Dimyristoylphosphatidylethanolamine on Cells that are Resistant to Methotrexate", <i>Biochimica et Biophysica Acta</i> , 885:129-135 (1986)			
	AY4	Kinsky <i>et al.</i> , "Circumvention of the Methotrexate Transport System by Methotrexate-Phosphatidylethanolamine Derivatives Effect of Fatty Acid Chain Length", <i>Biochimica et Biophysica Acta</i> , 921:96-103 (1987)			
↓	AZ4	Ko <i>et al.</i> , "Efficient Synthesis of Novel Carbocyclic Nucleosides via Sequential Claisen Rearrangement and Ring-Closing Metathesis", <i>Tetrahedron Letters</i> , 43:6399-6402 (2002)			
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/L.H./	AA5	Kofoed <i>et al.</i> , "Regiosomers of 2',3'-Dideoxynucleosides Related to 2-(Phosphonylmethoxy)ethyl Nucleosides", <i>Bulletin de la Societe Chimique de France</i> , 134:59-65 (1997)			
	AB5	Kraus, "New Phosphonate Analogues of 3'-thia-2',3'-Dideoxycytidine (BCH-189) Synthesis and Anti-HIV Evaluation", <i>Nucleosides & Nucleotides</i> , 12(2):157-162 (1993)			
	AC5	Leff <i>et al.</i> , "The Antidiabetic PPAR γ Ligands: An Update on Compounds in Development", <i>Curr. Med. Chem. - Immun., Endoc. & Metab. Agents</i> 2(1):33-47 (2002)			
	AD5	Lewandowicz <i>et al.</i> , "Achieving the Ultimate Physiological Goal in Transition State Analogue Inhibitors for Purine Nucleoside Phosphorylase", <i>The Journal of Biological Chemistry</i> , 278(34):31465-31468 (2003)			
	AE5	McGuigan <i>et al.</i> , "Synthesis and Anti-HIV Activity of Some Novel Chain-Extended Phosphoramidate Derivatives of d4T (Stavudine): Esterase Hydrolysis as a Rapid Predictive Test for Antiviral Potency", <i>Antiviral Chemistry & Chemotherapy</i> , 9:109-115 (1998)			
	AF5	McGuigan <i>et al.</i> , "Synthesis, Anti-Human Immunodeficiency Virus Activity and Esterase Lability of Some Novel Carboxylic Ester-Modified Phosphoramidate Derivatives of Stavudine (d4T)", <i>Antiviral Chemistry & Chemotherapy</i> , 9:473-479 (1998)			
	AG5	Mendes <i>et al.</i> , "Synthesis, Stability and In Vitro Dermal Evaluation of Aminocarbonyloxymethyl Esters as Prodrugs of Carboxylic Acid Agents", <i>Bioorganic & Medicinal Chemistry</i> , 10(3):809-816 (2002)			
	AH5	Menéndez-Arias, "Targeting HIV: Antiretroviral Therapy and Development of Drug Resistance", <i>TRENDS in Pharmacological Sciences</i> , 23(8):381-388 (2002)			
↓	AI5	Ono-Nita <i>et al.</i> , "Novel Nucleoside Analogue MCC-478 (LY582563) Is Effective Against Wild-Type or Lamivudine-Resistant Hepatitis B Virus", <i>Antimicrobial Agents and Chemotherapy</i> , 46(8):2602-2605 (2002)			
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/L.H./	AJ5	Pankiewicz <i>et al.</i> , "Novel Mycophenolic Adenine Bis(phosphonate) Analogues As Potential Differentiation Agents Against Human Leukemia", <i>J. Med. Chem.</i> , 45(3):703-712 (2002)	
	AK5	Parang <i>et al.</i> , "Novel Approaches for Designing 5'-O-Ester Prodrugs of 3'-Azido-2', 3'-Dideoxythymidine (AZT)", <i>Current Medicinal Chemistry</i> , 7(10):995-1039 (2000)	
	AL5	Pokrovskii <i>et al.</i> , "Comparative Analysis of HIV-1 Resistance to AZT and AZT H-Phosphonate in a Cell Culture", <i>Doklady Biochemistry and Biophysics</i> , 384:152-154 (2002)	
	AM5	Prashad <i>et al.</i> , "An Efficient and Large-Scale Enantioselective Synthesis of PNP405: A Purine Nucleoside Phosphorylase Inhibitor", <i>J. Org. Chem.</i> , 67(19):6612-6617 (2002)	
	AN5	Ray <i>et al.</i> , "Role of Purine Nucleoside Phosphorylase in Interactions between 2', 3'-Dideoxyinosine and Allopurinol, Ganciclovir, or Tenofovir", <i>Antimicrobial Agents and Chemotherapy</i> , 48(4):1089-1095 (2004)	
	AO5	Roberts, "Development of the Route to the New Anti-AIDS drug Abacavir: A highlight of Academic/Industry Liaison", <i>IDrugs</i> , 1(8):896-899 (1998)	
	AP5	Rosowsky <i>et al.</i> , "Methotrexate Analogues—27", <i>Biochemical Pharmacology</i> , 35(19):3327-3333 (1986)	
	AQ5	Rosowsky <i>et al.</i> , "Methotrexate Analogues, 32, Chain Extension, α -Carboxyl Deletion, and γ -Carboxyl Replacement by Sulfonate and Phosphonate: Effect on Enzyme Binding and Cell-Growth Inhibition", <i>J. Med. Chem.</i> , 31(7):1326-1331 (1988)	
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	AS5	Sauber <i>et al.</i> , "A New Esterase for the Cleavage of Pivalic Acid-Containing Prodrug Esters of Cephalosporins", <i>Enzyme and Microbial Technology</i> , 19:15-19 (1996)	
↓	AT5	Schultz, "Prodrugs of Biologically Active Phosphate Esters", <i>Bioorganic & Medicinal Chemistry</i> , 11:885-898 (2003)	
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		18477.031		10/740,694	
		APPLICANTS			
		Murty N. ARIMILLI <i>et al.</i>			
		FILING DATE		GROUP	
		December 22, 2003		1648	
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/L.H./	AU5	Sekiya <i>et al.</i> , "2-Amino-6-Arylthio-9-[2-(Phosphonomethoxy) Ethyl] Purine Bis(2,2,2-Trifluoroethyl) Esters as Novel HBV-Specific Antiviral Reagents", <i>Journal of Medicinal Chemistry</i> , 45(14):3138-3142 (2002)			
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	AW5	Siddiqui <i>et al.</i> , "Design and Synthesis of Lipophilic Phosphoramidate d4T-MP Prodrugs Expressing High Potency Against HIV in Cell Culture: Structural Determinants for in Vitro Activity and QSAR", <i>Journal of Medicinal Chemistry</i> , 42(20):4122-4128 (1999)			
	AX5	Sintchak <i>et al.</i> , "The Structure of Inosine 5'-Monophosphate Dehydrogenase and the Design of Novel Inhibitors" <i>Immunopharmacology</i> , 47:163-184 (2000)			
	AY5	Srinivas <i>et al.</i> , "Metabolism and <i>In Vitro</i> Antiretroviral Activities of Bis(Pivaloyloxymethyl) Prodrugs of Acyclic Nucleoside Phosphonates", <i>Antimicrobial Agents and Chemotherapy</i> , 37(10):2247-2250 (1993)			
	AZ5	Sturtz <i>et al.</i> , "Sur une nouvelle Approche de Pharmacomodulation du Methotrexate: Synthese D'analogues Gem-Diphosphoniques D'amethoptérine et de la N-10 Deaza Amethoptérine", <i>Medicinal Chemistry, C. R. Acad. Sci. Paris</i> , 10(2):739-742 (1990)			
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	AB6	Sturtz, <i>et al.</i> , "Synthesis of Gem-Bisphosphonic Methotrexate Conjugates and Their Biological Response Towards Walker's Osteosarcoma", <i>Eur. J. Med. Chem.</i> , 28:899-903 (1993)			
↓	AC6	Sturtz <i>et al.</i> , "A Study of the Delivery-Targeting Concept Applied to Antineoplastic Drugs Active on Human Osteosarcoma, I. Synthesis and Biological Activity in Nude Mice Carrying Human Osteosarcoma Xenografts of Gem-Bisphosphonic Methotrexate Analogues", <i>Eur. J. Med. Chem.</i> , 27(8):825-833 (1992)			
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/L.H./	AD6	Tan <i>et al.</i> , "Sequencing and Cloning of Human Prolylcarboxypeptidase (Angiotensinase C): Similarity to Both Serine Carboxypeptidase and Prolylendopeptidase Families", <i>The Journal of Biological Chemistry</i> , 268(22):16631-16638 (1993)			
↓	AE6	Valette <i>et al.</i> , "Decomposition Pathways and <i>in Vitro</i> HIV Inhibitory Effects of IsoddA Pronucleotides: Toward a Rational Approach for Intracellular Delivery of Nucleoside 5'-Monophosphates", <i>Journal of Medicinal Chemistry</i> , 39(10):1981-1990 (1996)			
	AF6	Vielhaber, "Bericht vom 3rd International Workshop on Salvage Therapy for HIV-Infection", <i>Deutsche Aids-Hilfe e.V. Fax Report zu HIV und AIDS</i> , Seite 4 (2000)			
	AG6	Waegell <i>et al.</i> , "A420983, a Novel, Small Molecule Inhibitor of LCK Prevents Allograft Rejection", <i>Transplantation Proceedings</i> , 34:1411-1417 (2002)			
	AH6	Wróblewski <i>et al.</i> , "Synthesis of (1R,2S)- and (1S,2S)-3-(4-Carbamoyl-1,2,3-triazol-1-yl)-1,2-Dihydroxypropylphosphonates, <i>Tetrahedron: Asymmetry</i> , 15:1457-1464 (2004)			
	AI6				
	AJ6				
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